



Publication number:

0 428 276 A2

1 2 1

EUROPEAN PATENT APPLICATION

- (2) Application number: 90311484.1
- $\frac{1}{2}$ Date of filing: 19.10.90

- © Int. C.E. **A01N 37/10,** A01N 37/02, A01N 39/00, A01N 31/04, A01N 43/30, A01N 31/14
- (8) Priority: 19.10.89 JP 272401/89
- Date of publication of application: 22.05.91 Bulletin 91/21
- Designated Contracting States: BE DE FR GB NL
- Applicant: Takasago International Corporation 19-22, Takanawa 3-chome Minato-ku Tokyo(JP)
- 2) Invenior: Sato, Toshiya, c/o Takasago Int. Corp. Kamata Div., 36-31, Kamata 5-chome Ohta-ku, Tokyo(JP) Inventor: Hata, Hamako, c/o Takasago Int. Corp. Kamata Div., 36-31, Kamata 5-chome Ohta-ku, Tokyo(JP)
- @ Representative: Moore, Anthony John et al Gee & Co. Chancery House Chancery Lane London WC2A 1QU(GB)

Acaricidal composition.

[☑] An acaricidal composition comprises, as active ingredient, one or more compounds selected from methyl
innamate, ethyl cinnamate, n-propyi cinnamate, isopropyi cinnamate, n-butyl cinnamate, isobutyl
innamate, in-hexyl cinnamate, allyl cinnamate, cinnamyl acottate, cinnamyl propionate, cinnamyl inoutyrate, cinnamyl isobutyrate, p-cresyl acetate, p-cresyl butyrate, p-cresyl isobutyrate, p-methylbenzyl propionate, β-phenoxyethyl alcohol, pnenoxyethyl acottate phenoxyethyl propionate, phenoxyethyl in-butyrate,
phenoxyethyl isobutyrate, methyl phenylacetate, ethyl phenylacetate, dibenzyl ether, heliotropin, methyl diphenyl
ether and 2-methyl-1-(methylbicyclo[2,2.1]hept-5-en-2-yl)-1-periten-3-oi.

ACARICIDAL COMPOSITION

This invention relates to an acaricidal composition which is free from harmful effects on human beings and is very effective for exterminating house dust acari.

House dust acari inhabit, and propagate mainly in, highly moist places, for example, on the surface of floors, under or within foor coverings such as tatami or carpet, or within bedclothes. Recently, Dermatophagoides including Dermatophagoides peronyssinus and Dermatophagoides farinae, which constitute 90% of house dust acari, have become a serious problem since they are important altergens causing bronchial asthma. 3 isorgic rhinitis and atopic dermatitis.

The most effective method for exterminating these acon is to ventilate and dry the house well. However, the recent increase in the number of houses having a closed structure and changes in itle style make it more and more difficult to ventilate a room sufficiently. Under these circumstances, the damage caused by acon has become more and more serious.

In order to exterminate these again, various agaricides (for example organophosphorus compounds such as fenitrothion, fenthion, dichloryos, diazinon; carbamate etimpounds such as propoxur, carbaryl; pyrethroid compounds such as resmethrin, phenothrin, permethrin have been applied in the form of 15 aemsol, fumigant, insecticidal sheet or impregnating agent for, e.g., carpets. Furthermore it was recently proposed to use compounds other than those cited above for exterminating acar. For example, JP-A-61-57501 discloses using a combination of adarticidal compounds such as benzy) benzoate, benzyl salicylate or dibityl phthalate with a powdery cleanser, and indicates that the adarcidal effect of benzyl benzoate has been physiologically particularly well studied. (The term "JP-A" as used herein means an "unexamined 20 published patent application".) JP-A-61-91103 discloses an acardole which comprises benzyl benzeate and an aliphatic hydrocarbon as the major components. Further, JP-A-61-97603 discloses benzyl salicytate and phenethyl benzoare, white JP-A-62-33106 discloses phenyl salicitate, phenyl benzoate id phenylamine. memyl á-naphthyl ketone and doumarin each as an active ingredient for an adaricide. Furthermore, JP-A-64-19004 discloses an adarticide comprising benzaldehyde or perillaldi-hyde. I-carvone or d-carvone, 25 mernyl salicylate or ethyl salicylate, or methyl benzoate or ethyl benzoate as an active ingredient. Regarding natural substances, furthermore, JP-A-63-104905 discloses that terpene compounds are available as scari-prevention agents. Furthermore, it is known that other vegetable assential bils (for example, bitter almond cil, wintergreen oil) show an acaricidal effect (F. Watanabe et al., Shoyal-ugaku Zasshi, 43 [2], 163-168 (1989)).

However, typical known acaricidal compounds (particularly organophosophorus compounds) generally show a high toxicity and exert undesirable effects on the human body. Therefore, it is undesirable to use these compounds in contined conditions or around houses. These compounds are further disadvantageous in that their effects on Dermatophagoides causing altergic diseases are limited. On the other hand, pyrethroid compounds are expensive and show only limited effects on house dust acari, though they are less toxic in general. Other acaricidal compounds are also disadvantageous in their limited effects on Dermatophagoides.

Accordingly, it has been urgently required to develop an acarbide which is very safe with respect to effects on the numan body, can be eastly used anywhere in the house, and yet exerts a powerful effect in exterminating a number or house dust alari, including Dermatophagoides, which cause altergoldissales.

We have found that the following compounds, which have been used as perfumes in focilis and cosmetics for a long time and have thus been proved to be harmless to human poings, are nightly effective in the extermination of house dust acari.

According to the present invention there is provided an acancidal composition comprising as the active ingradient one or more comprising selected from among methyl clinnamate, ethyl clinnamate. Propoglas clinnamate, isopropyl clinnamate, n-butyl crinnamate, isopropyl clinnamate, n-butyl crinnamate, isopropyl clinnamate, inversity crinnamate, allyl clinnamate, clinnamyl acetate, clinnamyl propionate, clinnamyl acetate, clinnamyl isobutyrate, p-cresyl acetate, p-cresyl butyrate, p-cresyl isobutyrate, p-methylbenzyl propionate, al-phenoxyethyl alcohol, phenoxyethyl acetate, phenoxyethyl propionate, phenoxyethyl inchipropionate, phenoxyethyl alcohol, phenylacetate, ethyl phenylacetate, dibenzy, ether, heliotropia, methyl diphenyl ether and 2-methylbogolog.2.1 [hept-5-en-2-yl)-1-penten-3-ol.

The acari to be exterminated with the acaricidal composition of the present invention include not only house dust acari inhabiting and propagating indoors, for example, Pyrogryphidae such as Dermatophagoides farinae and Dermatophagoides pteronyssinus; Aharidae such as Typophagus purrescentiae and Aleuroglyphus ovatus; Gly typnagidue such as Glycyphagus privatus and Glycyphagus domesticus; and Cheyletidae such as Cheyletius malacoansis and Cheyletia forus; but an mal-parastic acari, for example.

Marcronyssidae such as Ornithonyssus bacoti and Ormithonyssus sylviarum.

The acaricidal composition of the present invention may consist of one of the above-mentioned active ingredients or a combination thereof, as such. In general, however, it may be formulated into an pil pregaration, emuisifiable concentrare, wettable powder, spray, aerosol, furnigant, ocating, detergent, dust, grarules or capsules by supporting on a solid or liquid carrier and optionally adding various additives, for example, film-forming agent, emuls fier. Sticking agent, dispersant, wetting agent, stabilizer, propellant and willafility-controller, if required.

E-ambies of the solid carrier to be used herein include mineral powders such as silicic acid, Factin, as thered carbon, bentonite, diatomaceous earth, talc and calcium parbonate; vegetable powders such as wheat flour and starch; and synthetic polymer powder such as polyvinyl chloride powder. Examples of the highlicitation and starch; adiptatic hydrocarbone such as hoxine, kerosene and coal oil archaet hydrocarbone such as hoxine, kerosene and coal oil archaet in discretion such as benzere, tollene and kylene; halogenated hydrocarbons such as dishlordethane and circon totrachforide alcohols such as ethanol, isopropyl alcohol and ethylene glynot; ketones such as atenne, methyl ethyl ketone and cyclonoxianono; others such as tetrahydrofuran, dimetholyethane and dettyl ether, esters such as ethyl acetate; nitriles such as aceton; rile; socid amides such as dimethylformamics; and vegetable oils such as skybean oil and cotton seed oil.

Stamples if the filtriforming agent include cellulose derivatives vinys resins, ally direstins, unchanne resins, sitioper resins, acry to resins, or brinders and puly inyl acount. Examples of this emulsifier, sticking agent and dispersant include surfactants such as places polyoxyethyltino allylary: ethers, polyoxyethylene fatty acid esters, fatty acid glycerols, somiran tatty acid esters, higher alcohol suffacts, and alky arylstufenic acid safts. Examples of the propollant include is useful petroleum gap. From gas and dimethylillether. Examples of the volatifity-controller include it by locecane and tyclichoid cane.

Furthermore, the active ingrenerit(s) may be used together with publimating insecticides such an user distribution properties in aprilhalene or camphor so as to give a sublimating solid preparation.

Moreover, the spanoisal composition of the present invention may contain, for example, various some attornal insecticides, accidedes, synergists, harmful insect repellents, redent repellents, hacterialdes perfumes or colorants used for exterminating harmful insects, such as fentrothion, proposuli or especificing.

The content of the abore-mentioned active ingredient in the aparcidal composition of the present content may vary depending on the formulation, application means and the place to be expired in the presently preferable that the rotal content of the active ingredient(s) ranges from 0.1 to 50% by regignt (in the use of the wettable powder or emulsitable concentrate) and from 0.1 to 30% by weight (in the use of of place attion or series), respectively

The acandidal composition of the present invention thus prepared may be applied to, for example, there, taram, carpets, bedo others, sofas pillows on closests by depositing spraying, coating, transpring or pacement. Attendatively, it may be used as a deterged for human or bot animals. The dose in preferably, approximately 80 mg or more per m³ of the area to be treated or approximately 8 mg or more per m³ of the scale to he treated, in terms of the rotal amount of the active ingredient.

in addition to the above trimulations, the acarcide of the present invention may be formulated into tiem orient or enstructional material having an acardidal activity by supporting the active impreciants on an expreportian substrate. Examples of the substrate to be used herein include sheets of synthetic resins such as equip-hyllone, propriory-inc, hylline, potyvinyl chloride or polyesters, animal or vegetable from an area at some inorganic fibrous materials or inorganic fibrous materials and animal, vegetable or inorganic fibros; mixed tables or non-worsen fathers, foils or films of merals such as aluminum, stainless steel or ainc; faminaries of the above-mentioned sheets and various nutural wooden materials and plastics molded articles employed for constructional outposes. The active ingredient of the acardidal composition of the present mention is operated in these substrates by loading, impregnating, depositing or carabricating to give an acardidal or raterial. The amount of the active ingredient in the substrate is not particularly restricted out may be optionally selected in the case of impregnation, it is preferable to use the active ingredient in the saturation.

The adamoidal material thus obtained may be prefirably used, for example, in the following manner: A pullymer sheet (for example, notyprubylene) impregnated with the active ingredient of the present invention is placed under *tatami*, carpets or sofas. In this case, it is preferable to use the active ingredient at a ratio of firm approximately 0.5 to 20 giger unit area. The impregnation of the polymer with the active ingredient makes the sustained release of the active ingredient possible, which brings about a sustained acarroidal effect.

The effects of the active ingredients of the present invention were examined by using Dermatophagoides pteronyssinus, which is one of Dermatophagoides and is generally less sensitive to chemicals, by the following procedure.

Namely, a filter paper (6 mm x 5 mm) is impregnated with each test compound in such a manner as to 5 give the definite concentration. A liquid compound is used as such while a solid one is dissolved in accordance with a method reported by Watanaoe et al., Shoyakugaku Zasshi, 43 [2], 163-168 (1989) the filter sheet is introduced into a cylindrical container (approximately 20 cc) containing 50 to 80 head of Dermatophagoides pteronyssinus together with a bait. The container is then sealed with a Tellon stopper and allowed to stand in an incubator at 25° C. After 24 hours and 48 hours, the life or death of the aceri is avamined under a stereoscopic microscope or a loupe (x 25) and evaluated. The procedure is repeated thrice and the lethality is calculated according to the following equation. Table 1 shows average values. Lethality (%) = (X + Y)X × 100

X: number of living acari in untreated plot; and

Yi number of living acari in treated plot.

In Table 1, a mixture of Test Compounds is expresed by the Compound Number of each component.

For comparison, permethrin and benzyl saficylate, which are conventional acaricides, were also exclusted in the same manner. The results are shown in Table 1.

:5

30

35

active (0.04 g/m ²) After 48 hours	100	100	06	86	06	75	100	62	100	82	68	72	8.5	86
Dose of ingredient After 24 hours	100	100	7.8	8 2	7.5	7.1	76	5.8	72	53	9	53	6.2	80
active (0.08 q/m2) After 48 hours	100	100	100	100	100	100	100	100	100	100	100	100	100	100
Dose of ingredient After 24 hours	100	100	98	92	9.0	8.2	88	8.2	79	100	100	8.0	100	8.5
Test Compound (blending ratio)	Methyl cinnamate	Ethyl cinnamate	n-Propyl cinnamate	Isopropyl cinnamate	n-Butyl cinnamate	Isobutyl cinnamate	Isoamyl cinnamate	n-Hexyl cinnamate	Allyl cinnamate	Cinnamyl acetate	Cinnamyl propionate	Cinnamyl n-butyrate	Cinnamyl isobutyrate	p-Cresyl acetate
Compound No.	1)	(2)	(3)	(4)	(5)	(9)	(7)	(8)	(6)	(10)	(11)	(12)	(13)	(14)
Type	Single compound													
	Compound Test Compound After After After After No. (blending ratio) 24 hours 24 hours	Compound Test Compound After 180. [1] Methyl cinnamate 100 100	Compound Test Compound Afrer Afrer Afrer No. (blending ratio) 24 hours 48 hours (1) Methyl cinnamate 100 100 (2) Ethyl cinnamate 100 100	Compound Test Compound Ingredient (0.08 g/m²) After After After After After After After (1) Methyl cinnamate 100 100 100 (2) Ethyl cinnamate 86 100 100	Compound Test Compound After (0.08 q/m²)	Compound Test Compound Ingredient (0.08 q/m2)	Compound Test Compound Ingredient (0.08 g/m²) After After	Compound Test Compound Ingredient (0.08 q/m²)	Compound Test Compound Ingredient Co.08 q/m²	Compound Test Compound Ingredient 0.08 q/m²; 1	Compound Test Compound Ingredient Co.08 q/m2) Affer Co.08 q/m2 Co.08 q			

				EP	0 42	8 276	6 A2							
	active (0.04 g/m²) After 48 hours	76	85	96	100	86	85	86	100	100	100	100	100	100
ity	Dose of ingredient After 24 hours	83	7.8	86	96	986	7.8	8.2	100	100	100	100	100	100
Lethallty	active (0.08 q/m²) After 48 hours	100	160	100	100	100	100	100	100	100	100	100	100	100
0.00	Dose of ingredient After 24 hours	92	06	36	100	100	100	100	100	100	100	100	100	100
	Test Compound in (blending ratio)		p-Cresyl isobutyrate	p-Methylbenzyl propionate	B-Phenoxyethyl alcohol	Phenoxyethyl acetate	Fhendayethyl propionate	Phenoxyethyl n-butyrate	Phenoxyethyl isobutyrate	Methyl phenylacetate	Ethyl phenylacetate	Dibenzyl ether	Heliotropin	Methyl diphenyl ether
	Compound No.	(15)	(16)	(17)	(18)	(19)	(50)	(21)	(22)	(23)	(24)	(25)	(26)	(27)
	Type	Single compound												

4()

				2P V 4	28 21	'0 A2							
	Dose of active ingredient (0.04 g/m²) After After 24 hours 48 hours	100	100	100	100	100	100	100	100	100	100	100	100
itv	Dose of ingredient After 24 hours	69	100	100	100	100	86	06	100	100	5.0	100	90
Lethality	active (0.08 g/m²) After 48 hours	100	100	100	100	100	100	100	100	100	100	100	100
ont'd)	Dose of active ingredient (0.08 g/m²) After After 24 hours 48 hours	100	100	100	100	100	100	100	100	100	106	100	100
TABLE 1 (cont'd)	wound Test Compound (N)	2 Methyl-1-(methylbi- cyclo[2.2.1]hcpt-5-en- 2-yl)-1-penten-3-ol	(2)/(15) (1/1)	(2)/(18) (1/1)	(2)/(24)/(4/1)	(2)/(27) (1/1)	(14)/(18) (1/1)	(14)/(23) (1/1)	(14)/(25) (1/1)	(14)/(27) (1/1)	(2)/(18)/(24)/(1/1/1)	(1/1/1) (22)/(32)/(54)	(25)/(26)/(27) (1/1/1)
	Сощення Вуре Н5.	Single compound (28)	Mixed Composition										

TABLE 1 (cont'd)

a()

	active (0.04 g/m ²)	After	48 hours	83	32
ity	Dose of active ingredient (0.04 g/m²)	After	24 hours	63	25
Lathality	Dose of active	After	48 hours	0.6	7.0
	Dose of	After	24 hours	82	50
		Test Compound	(blending ratio)	hrin	Benzyl salicylate
		Compound	NC.	Permethrin	Benzyl
			Type	Comparison	

As Table 1 clearly shows, the active ingredients of the acaricide of the present invention were superior to permethrin and benzyl salicylate for exterminating Dermatophagoides pteronyssinus.

To further idustrate the present invention, and not by way of limitation, the following Examples will be

5 given.

EXAMPLE 1

10

15

Oil Preparation: (parts by weight) Ethyl cinnamate 2 Isopropyl alcohol 98 Total 100

20

The above components were mixed under stirring to give a homogeneous oil preparation.

25

EKAMPLE 2

30

35

Emu sifiable Concentrate	Э
	rparts by weight)
Cinnamyl acetate	20
Sorbitan monostearate	:0
≯ylene	70
Total	100

10

The above components were mixed under stirring to give a homogeneous emulsion.

EXAMPLE 3

50

15

Dust:	
	(parts by weight)
3-Phenoxyethyl alcohol	10
Silicic anhydride	5
Talc	85
Total	100

The above components were intimately mixed to give a homogeneous dust.

EXAMPLE 4

Dust:

(parts by weight)

Methyl phenyla: etate 40
Soft polyvinyl chloride powder 60

Total 100

The above components were stirred at reom temperature over day and night to allow the polyvinyl chloride powder to absorb the methyl phenylac-tate. Thus a dust was prepared.

EXAMPLE 5

2.5

317

35

40

1-)

15

Detergent:	
	(parts by weight)
p-Gresyl butyrate Polyoxyethylene nanylphenyl ether water	10 25 65
Total	100

The above components were intimately mixed to give a homogeneous detergent.

E · AMPLE 6

45

Aerosol:	
	(parts by weight)
Ethyl phenylacetate	10
Dimethoxyethane	40
Liquefied petroleum gas	50
Tota-	100

55

The ethyl phenylacetate and dimethoxyethane were mixed under stirring and then introduced into an aerosol container. After providing a valve, the liquefied petroleum gas was fed thereinto through the valve under a pressure to give an aerosol.

EYAMPLE 7

5	Aerosol:	
		(parts by weight)
	p-Cresyl butyrate	5
10	Methyl diphenyl ether	5
	Xylene	10
	Illuminating kerosene	30
	Liquefied petroleum gas/dimethyl ether mixture (ratio by volume = 1:1)	50
15	Total	100

The above components except the mixture of liquefied petroleum gas and dimethyl ether were mixed under stirring and then introduced into an aerosol container. After providing a valve, the mixture of liquefied petroleum gas and dimethyl ether was fed thercinto through the valve under a pressure to give an aerosol.

E: AMPLE 8

Sheet material:	
	(parts by weight)
Methyl pher ylacetate	20
Ethyl cellulc-se	10
Ethanol	70
Total	100

The above components were mixed under stirring, and a polyethylene pulp non-woven fabric was impregnated therewith in such a manner as to give a ratio of methyl pheny acetate of 1 g/m². Thus a sheet material was obtained.

EKAMPLE 9

45

	Sheet material:				
		(parts by weight)			
	Ethyl cinnamate	10			
	Dipenzyl ether	10			
	Ethyl cellulose	10			
	Ethanol	70			
ĺ	Total	100			

The above components were mixed under stirring, and a polyothylene pulp non-woven fabric was impregnated therewith in such a manner as to give a total amount of ethyl cinnamate and dibenzyl ether of 1 g/m². Thus a sheet material was obtained.

The acaricidal composition of the present invention exhibits an excellent effect of extermination house dust acari. Further, it is highly safe to human body and can be easily applied in the house, which makes it extremely advantageous.

10 Claims

- 1. An acarcidal composition comprising one or more compounds selected from methyl cinnamate, ethyl cinnamate, n-propyl cinnamate, isopropyl cinnamate, n-butyl cinnamate, isobutyl cinnamate, isopropyl cinnamate, isopropyl cinnamate, allyl cinnamate, cinnamyl acetate, cinnamyl propionate, cinnamyl n-butyrate,
- icinnamyl isobutyrate, p-cresyl acetate, p-cresyl butyrate, p-cresyl isobutyrate, p-methyloenzyl propionate, g-phenoxyethyl alcohol, phenoxyethyl acetate, phenoxyethyl propionate, phenoxyethyl n-butyrate, phenoxyethyl isobutyrate, metnyl phenyl-acetate, ethyl phenylacetate, dibenzyl ether, heliotropin, methyl diptienyl ether and 2-methyl-t-(methylbcyclo(2.2.1)heot-5-en-2-y)-1-penten-3-o- as an active ingredient.
 - 2. An acaricidal composition as claimed in claim 1 and containing a solid or liquid carrier.
- 3. An acaricidal composition as claimed in claim 2, wherein the active ingredient is present in an amount of from 0.1 to 50% by weight.
 - 4. An acarrora, composition as claimed in claim 3, wherein said composition is in the form of a wettable powder or an emulsifiable concentrate.
- 5. An acaricidal composition as claimed in claim 2, wherein the active ingredient is present in an amount of irom 0.1 to 30% by weight.
 - An acaricidal composition as claimed in claim 5, wherein said composition is in the form of an oil preparation or an aerosol.
- 7. A method of exterminating house dust acarr, which comprises applying an acaricidal composition comprising, as active ingredient, one or more compounds selected from methyl cinnamate, electronic comprising, as active ingredient, one or more compounds selected from methyl cinnamate, isoamyl cinnamate, isoamyl cinnamate, isoamyl cinnamate, isoamyl cinnamate, isoamyl cinnamate, cinnamyl isoamyl in-butyrate, cinnamyl isoamyl propionate, cinnamyl in-butyrate, cinnamyl isoamyl acottal, percesyl actate, percesyl isoamyl propionate, percetyl propionate, a-phenoxyethyl nebuty acte, phenoxyethyl propionate, phenoxyethyl in-buty acte, phenoxyethyl percetyl phenoxyethyl contamate, inches phenoxyethyl phenoxy
 - 8. A method as claimed in claim 7, wherein said house dust acari are Dermatophagoides.
 - 9. Use as an acarcide of any of the compounds listed in claim 1, either singly or in any combination thereof



Europaisches Patentamt European Patent Office Office européen des brevets Publication number:

0 428 276 A3

EUROPEAN PATENT APPLICATION

- Application number 90311484.1
- Date of filing: 19.10.90

© Int. Cl.5 A01N 37/10, A01N 37/02, A01N 39/00, A01N 31/04, A01N 43/30, A01N 31/14

- Priority: 19.10.89 JP 272401/89
- Date of publication of application: 22.05.91 Bulletin 91/21
- Designated Contracting States: BE DE FR GB NL
- Oate of deferred publication of the search report. 14.07.93 Bulletin 93/28
- Applicant: Takasago International Corporation
 19-22, Takanawa 3-chome Minato-ku
 Tokyo(JP)
- (*) Inventor Sato, Toshiya, c/o Takasago Int. Corp.
 Kamata Div., 36-31, Kamata 5-chome
 Ohta-ku, Tokyo(JP)
 Inventor: Hata, Hamako, c/o Takasago Int.
 Corp.
 Kamata Div., 36-31, Kamata 5-chome
 Ohta-ku, Tokyo(JP)
- Representative: Moore, Anthony John et al Gee & Co. Chancery House Chancery Lane London WC2A 1QU (GB)

Acaricidal composition.

§ 1 An acaricidal composition comprises as active ingredient, one or more compounds selected from methyl cinnamate, ethyl cinnamate, in-propyl cinnamate, isopropyl cinnamate, in-butyl cinnamatic, isomyl cinnamate, isomyl cinnamate, isomyl cinnamate, isomyl cinnamate, cinnamyl propionate, allyl cinnamyl in-butyrate, cinnamyl isotrutyrate, cinnamyl isotrutyrate, p-cresyl acetate p-cresyl butyrate, p-cresyl isobutyrate, p-mothylbonzyl propionate, p-penoxyethyl alcohol, phenoxyethyl acetate, phenoxyethyl isobutyrate, methyl phenylacetate, athyl phenylacetate dibenzyl ether, heliotropin, methyl cinhenyl ether and 2-methyl-1-(methylbicyclo[2.2.1]-hept-5-en-2-yl)-1-penten-3-ci.

EP 90 31 1484

Category	Citation of document with it of relevant pa	edication, where appropriate,	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)	
X	CHEMICAL ABSTRACTS, 16th January 1978, no. 17260z, Columbu DESHPANDE et al.: " activity of Ocimum PESTICIDES 1977, 11 * Abstract *	vol. 88, no. 3, page 163, abstract s, Ohio, US; R.S. Insecticidal basilicum Linn", &	1	A 01 N 37/10 A 01 N 37/02 A 01 N 39/00 A 01 N 31/04 A 01 N 43/30 A 01 N 31/14	
X	US-A-3 259 648 (H. * Whole document *	E. HENNIS)	1-9		
A	EP-A-0 235 722 (BA	SF AG)			
X	FR-A-2 392 602 (BL * Page 1, lines 1-3 15-19; tables, clai		1-9		
X	US-A-4 368 207 (BL * Column 1, line 58 18; tables; claims		1-9		
Ρ,Χ	WO-A-8 912 673 (VA * Page 5, lines 6-1 1-10; page 22, line		1-9	TECHNICAL FIELDS SEARCHED (Int. CL5)	
X	CHEMICAL ABSTRACTS, 7th May 1979, page 147035g, Columbus, 661 (YOZMETIKAI ES VALLALAT) 28-11-197 * Abstract *	169, abstract no. Ohio, US; & HU-A-15 HAZTARTASVEGYIPARI	1-9		
	The process course report has	seen trans up for all claims			
Place of GENTS THE HAGUE		Date of completion of the sewich $0.1-0.2-1.991$	DON	DONOVAN T.M.	
X : pa Y : pa do A - ter	CATEGORY OF CITED DOCUME ricularly relevant if taken alone ricularly relevant if combined with an cument of the same category inhological background in myrice discissive	NTS T theory or prince E rearlier patent of after the fuling other D document cired L document cired	ple underlying th ocument, but put date in the applicatio for other reasons	e invention dished on, or	



CL	CLAIMS INCURRING FEES					
The present	European parent application comprised at the time of hising more than fen claims.					
	All claims lees have been paid within the prescribed time limit. The present European search report has been drawn up for all claims.					
	Only part of the claims fees have been paid within the prescribed lime limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been daid.					
	namely claims.					
	No claims fees have been paid within the prescribed time limit. The present European search report has been grawn up for the Arst ten claims.					
	CK OF UNITY OF INVENTION					
	Division considers that the present European patent application does not comply with the requirement of unity of idrelates to several inventions or groups of inventions.					
namely:	o relates to several inventions of Groups or Inventions;					
9	ee sheet -B-					
٥	ee sheet -B-					
	All further search less have been paid within the fixed time limit. The present European search report has been drawn up for all claims.					
M	Only part of the further search fees have been paid within the fixed time, irrid, The present European search report has been drawn up for those parts of the European palent application which relate to the inventions in respect of which search fees have been paid.					
	namely claims: points 1.,3.,4.,5., and 6.					
	None of the further search fees has been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which retate to the Invention first mentions of inhe claims.					
	namely claims					

EUROPEAN SEARCH REPORT

2

EP 90 31 1484

	Citation of document with ind	ERED TO BE RELEVA	Relevant	CLASSIFICATION OF THE
ategory	of relevant pass		to claim	APPLICATION (Int. Cl.5)
X	DERWENT CENTRAL PATE ABSTRACTS JOURNAL, so week Ell, 12th May 1 abstract no. 21087E/ Publications Ltd, Lo 024 303 (MITSUI TOAT 08-02-1982	ection C: AGDOC, 982, class CO3, 11, Derwent ndon, G8; & JP-A-57	1-9	
x	FR-A- 674 743 (IG * Whole document *	FARBENINDUSTRIE AG)	1-9	
X	"The Merck Index", e Merck & Co., Inc., R * Page 1078, compound	ahway, NJ, US	1-9	
P,X	WO-A-9 009 738 (CHA PRODUCTS LTD) * Whole document *	RWELL CONSUMER	1-9	
				TECHNICAL FIELDS SEARCHED (Int. CL5)
ļ				
į				
			+	
				1
				1
	The present course report how bee	•		
THE	Place of Geneta HAGUE	Date of completion of the search 01-02-1991	DONG	Examer DVAN T.M.
X : part	CATEGORY OF CITED DOCUMEN I ticularly relevant if taken alone ticularly relevant if combined with anoth ument of the same category, anological background	E earlier paten after the filli er D : document cr	nciple underlying Thi t document, but pub- ing date ted in the application led for other reasons	lished on, or

European Patent
Office

EP 90 31 1484 -B-

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application loces not comply with the requirement of unity of invention and relates to several inventions or groups of inventions.

namety:

- 1. Claims 1-9 (partially), as far as the active ingredient is one or more of methyl cirnamate, sthyl cirnamate, inspondy. Cirnamate, inspondyl cirnamate, insolved cirnamate, isodoutyl cirnamate, isodoutyl cirnamate, isodoutyl cirnamate, cirnamyl acetate, cirnamyl propionate, cirnamyl acetate cirnamyl propionate, cirnamyl nebutyrate or cirnamyl isodotyrate, optimally with one or more of proresyl acetate, proresyl but, nate, propesyl isodotyrate, promethylacetate, propionate, parhendryethyl ilochol, phendryethyl acetate, phendryethyl propionate, phendryethyl industryate, phendry ethyl isobotyl ste, methyl phendracetate, ethyl phenylocetate, dicencyl ether, neligorogin, methyl dippenyl ether or 2-methyl-1- methylbicyclo(2.2.1)hept-5-en-1-v1)- 1-penton-Dool.
- 2. Claims 1-9 (partially), as far as the active ingredient is one or more of precess' acetate, precess' buttrate, precess' isobut.rate or precipital pr
- 7. Claims 1-9 (partially), as far his the active ingredient is one or more of dephenosympholalcohol, thenosymphyl active, phenosympholalcohol, rebutyrate or phenosympholalcohol, rebutyrate or phenosymphyl isobutyrate, optionally with one or more of methyl phenylicethie, ethyl phenylacethie, disembyl ether, heliotropin, methyl diphenyl ether, then the phenylicethie, ethyl phenylacethie, of phenylicethie, bicyclo(2.1.))hept=5-en-2-yl)-1-panten-5-ol.
- 4. Claims 1-9 (partially), as far as the active ingredient is one or more of methyl phenylacetate or sthyl phenylacetate, optionally with one or more of dibersyl ether, heliotropin, methyl diprenyl ether or 2-methyl-1-(methy:bicyclo(2.2.1)hept-5-en-2-yl)-1-penten-T-ol.
- Claims 1-7 (partially), as far at the active ingredient is one or more of dibencyl other or metryl dichemyl ather, optionally with one or more of helictropin or Demetry: -1m(bicyclo(2.2.1) hept-5-en-2-yl)-1-perten-7-ol.
- 5. Claims 1-9 (partially), as far as the active ingredient is heliotropin, optionally with 2-methyl-1-(mgthylbicyclo-(2.2.1 hepr-6-en-2-yl)-1-penten-3-ol.
- Claims 1-9 (partially), as far %s the active ingredient is 2-methyl-1-(methylbicyclo(2.2.1)bept-5-am-2-yl)-1-penten-1-ol